ABSTRACT

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2.4-DISUBSTITUTED TRIAZINE DERIVATIVES

This invention concerns the use of the compounds of formula

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the *N*-oxides, the pharmaceutically acceptable addition salts, quaternary amines and the stereochemically isomeric forms thereof, wherein $-a^1=a^2-a^3=a^4$ - forms a phenyl, pyridinyl, pyrimidinyl, pyridazinyl or pyrazinyl with the attached vinyl group; n is 0 to 4; and where possible 5; R^1 is hydrogen, aryl, formyl, C_{1-6} alkylcarbonyl, C_{1-6} alkyl, C_{1-6} alkyloxycarbonyl or substituted C_{1-6} alkyl; each R^2 independently is hydroxy, halo, optionally substituted C_{1-6} alkyl, C_{2-6} alkenyl or C_{2-6} alkynyl, C_{3-7} cycloalkyl, C_{1-6} alkyloxy, C_{1-6} alkyloxycarbonyl, carboxyl, cyano, nitro, amino, mono- or di(C_{1-6} alkyl)amino, polyhalomethyl, polyhalomethyloxy, polyhalomethylthio, $-S(=O)_pR^4$, $-NH-S(=O)_pR^4$, $-C(=O)R^4$, -NHC(=O)H, $-C(=O)NHNH_2$, $-NHC(=O)R^4$, $-C(=NH)R^4$ or a 5-membered heterocyclic ring; p is 1 or 2; L is optionally substituted C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl or C_{3-7} cycloalkyl; or L is $-X-R^3$ wherein R^3 is optionally substituted phenyl, pyridinyl, pyrimidinyl, pyrazinyl or pyridazinyl; X is $-NR^1$ -, -NH-NH-, -N=N-, -O-, -C(=O)-, -CHOH-, -S-, -S(=O)- or $-S(=O)_2$ -; aryl is optionally substituted phenyl; for the manufacture of a medicine for the treatment of subjects suffering from HIV (Human Immunodeficiency Virus) infection.